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=> s lysine(w)rich and seed and protein and zein
L1 4 LYSINE(W) RICH AND SEED AND PROTEIN AND ZEIN

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L1 ANSWER 1 OF 4 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2004) on STN

ACCESSION NUMBER:

1998:49315 AGRICOLA

DOCUMENT NUMBER:

IND21243126

TITLE:

Lysine - ***rich*** modified gamma-

zeins accumulate in ***protein*** bodies

of transiently transformed maize endosperms.

AUTHOR(S):

Torrent, M.; Alvarez, I.; Geli, M.I.; Dalcol, I.;

Ludevid, D.

AVAILABILITY:

DNAL (QK710.P62)

SOURCE:

Plant molecular biology, May 1997. Vol. 34, No. 1. p.

139-149

Publisher: Dordrecht: Kluwer Academic Publishers.

CODEN: PMBIDB; ISSN: 0167-4412

NOTE:

Includes references

PUB. COUNTRY:

Netherlands

DOCUMENT TYPE:

Article

FILE SEGMENT:

Non-U.S. Imprint other than FAO

LANGUAGE:

English

AB During maize ***seed*** development, endosperm cells synthesize large amounts of storage ***proteins*** , alpha-, beta-, and gamma-

zeins , which accumulate within endoplasmic reticulum (ER)-derived
protein bodies. The absence of lysine in all ***zein***

polypeptides results in an imbalance in the amino acid composition of
maize ***seeds*** . We modified the maize gamma- ***zein*** gene
through the introduction of ***lysine*** - ***rich*** (Pro-Lys)n
coding sequences at different sites of the gamma- ***zein*** coding
sequence. Maize endosperms were transiently transformed by biolistic
bombardment with Lys-rich gamma- ***zein*** constructs under the

were observed. In contrast, when (Pro-Lys)n, sequences were inserted five residues from the C-terminal, the transcript was present but modified ***protein*** was not detected. These results suggest that only an appropriate positioning of Lys-rich inserts leads to the modified molecule displaying correct folding and stability. Subcellular localization analyses and immunoelectron microscopy studies on isolated ***protein*** bodies demonstrated that modified gamma- ***zeins*** accumulate within these organelles and co-localized with endogenous alpha- and gamma- ***zeins***. The studies reported here show the feasibility of manipulating the gamma- ***zeins*** gene in order to obtain stable and correctly targeted Lys-rich ***zeins*** in maize ***seeds***.

L1 ANSWER 2 OF 4 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN ACCESSION NUMBER: 1997:320606 BIOSIS

DOCUMENT NUMBER:

PREV199799611094

TITLE:

Lysine - ***rich*** modified gamma-

zeins accumulate in ***protein*** bodies of

transiently transformed maize endosperms.

AUTHOR(S):

Torrent, Margarita; Alvarez, Inaki; Geli, M. Isabel;

Dalcol, Ionara; Ludevid, Dolors [Reprint author]

CORPORATE SOURCE:

Dep. de Genetia Molecular, Centre d'Investigacio i

Desenvolupament, 08034 Barcelona, Spain

SOURCE:

Plant Molecular Biology, (1997) Vol. 34, No. 1, pp.

139-149.

CODEN: PMBIDB. ISSN: 0167-4412.

DOCUMENT TYPE:

Article

LANGUAGE:

English

ENTRY DATE:

Entered STN: 26 Jul 1997

Last Updated on STN: 26 Jul 1997

protein bodies. The absence of lysine in all ***zein***
polypeptides results in an imbalance in the amino acid composition of
maize ***seeds***. We modified the maize gamma- ***zein*** gene
through the introduction of ***lysine*** - ***rich*** (Pro-Lys),
coding sequences at different sites of the gamma- ***zein*** coding
sequence. Maize endosperms were transiently transformed by biolistic
bombardment with Lys-rich gamma- ***zein*** constructs under the
control of the 1.7 kb gamma- ***zein*** ***seed*** -specific
promoter and the cauliflower mosaic virus (CaMV) 35S promoter. When
(Pro-Lys), sequences were inserted contiguous to or in substitution of the
Pro-Xaa region of the gamma- ***zein***, high levels of ***protein***
were observed. In contrast, when (Pro-Lys)-n sequences were inserted five
residues from the C-terminal, the transcript was present but modified

protein was not detected. These results suggest that only an appropriate positioning of Lys-rich inserts leads to the modified molecule displaying correct folding and stability. Subcellular localization analyses and immunoelectron microscopy studies on isolated ***protein*** bodies demonstrated that modified gamma- ***zeins*** accumulate within these organelles and co-localized with endogenous alpha- and gamma-

zeins . The studies reported here show the feasibility of manipulating the gamma- ***zein*** gene in order to obtain stable and

1977:307431 CAPLUS ACCEDBLON NOMBEK: DOCUMENT NUMBER: 127:92732 TITLE: ***Lysine*** - ***rich*** modified .gamma.-***zeins*** accumulate in ***protein*** bodies of transiently transformed maize endosperms Torrent, Margarita; Alvarez, Inaki; Geli, M. Isabel; AUTHOR(S): Dalcol, Ionara; Ludevid, Dolors CORPORATE SOURCE: Departament de Genetica Molecular, Centre d'Investigacio i Desenvolupament, (CSIC), Barcelona, 08034, Spain SOURCE: Plant Molecular Biology (1997), 34(1), 139-149 CODEN: PMBIDB; ISSN: 0167-4412 PUBLISHER: Kluwer DOCUMENT TYPE: Journal LANGUAGE: English During maize ***seed*** development, endosperm cells synthesize large amts. of storage ***proteins*** , .alpha.-, .beta.-, and .gamma.-***zeins*** , which accumulate within endoplasmic reticulum (ER)-derived ***protein*** bodies. The absence of lysine in all ***zein*** polypeptides results in an imbalance in the amino acid compn. of maize ***seeds*** . We modified the maize .gamma.- ***zein*** ***lysine*** - ***rich*** (Pro-Lys)n coding the introduction of sequences at different sites of the .gamma. - ***zein*** coding sequence. Maize endosperms were transiently transformed by biolistic bombardment with Lys-rich .gamma.- ***zein*** constructs under the control of the 1.7 kb .gamma.- ***zein*** ***seed*** -specific promoter and the cauliflower mosaic virus (CaMV) 35S promoter. When (Pro-Lys)n sequences were inserted contiguous to or in substitution of the Pro-Xaa region of the .gamma.- ***zein*** , high levels of ***protein*** were obsd. In contrast, when (Pro-Lys)n sequences were inserted five residues from the C-terminal, the transcript was present but modified ***protein*** was not detected. These results suggest that only an appropriate positioning of Lys-rich inserts leads to the modified mol. displaying correct folding and stability. Subcellular localization analyses and immunoelectron microscopy studies on isolated ***protein*** bodies demonstrated that modified .gamma.- ***zeins*** accumulate within these organelles and co-localized with endogenous .alpha.- and .gamma.~ ***zeins*** . The studies reported here show the feasibility of manipulating the .gamma.- ***zein*** gene in order to obtain stable and correctly targeted Lys-rich ***zeins*** in maize ***seeds*** . ANSWER 4 OF 4 CAPLUS COPYRIGHT 2004 ACS on STN ACCESSION NUMBER: 1997:60561 CAPLUS DOCUMENT NUMBER: 126:87158 TITLE: Characterization of the variability in lysine content for normal and opaque2 maize endosperm AUTHOR(S): Moro, Gloverson L.; Habben, Jeffrey E.; Hamaker, Bruce

CORPORATE SOURCE:

Dep. Plant Sciences, Univ. Arizona, Tucson, AZ, 85721,

USA

SOURCE: Crop Science (1996), 36(6), 1651-1659

R.; Larkins, Brian A.

CODEN: CRPSAY; ISSN: 0011-183X

PUBLISHER: Crop Science Society of America, Inc.

DOCUMENT TYPE: Journal

increasing the content of this essential amino acid in endosperm ***proteins*** depends on understanding the mechanisms regulating the synthesis and accumulation of ***lysine*** - ***rich*** ***proteins*** . The variability for lysine and ***protein*** contents was studied in maize endosperm. Amts. of total ***protein***
, ***zeins*** , and non- ***zeins*** measured by microKjeldahl, and lysine content, estd. by amino acid anal., were detd. for 93 maize inbreds. Addnl., an ELISA was used to est. the relative content of the ***protein*** synthesis factor EF-1.alpha. in 20 selected genotypes. Considerable differences in lysine content were obsd. among normal and opaque2 genotypes, with the effect of the mutation being highly dependent on the genetic background. A high correlation was detected between the lysine content and the concn. of total non- ***zein*** ***proteins*** and EF-1.alpha.. An assay for EF-1.alpha. concn. may provide a simple and inexpensive method from breeding programs to select for improved ***protein*** quality. REFERENCE COUNT: 44 THERE ARE 44 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT => s seed and protein and gamma and zein 91 SEED AND PROTEIN AND GAMMA AND ZEIN => duplicate remove 12 DUPLICATE PREFERENCE IS 'AGRICOLA, BIOSIS, EMBASE, CAPLUS' KEEP DUPLICATES FROM MORE THAN ONE FILE? Y/(N):n PROCESSING COMPLETED FOR L2 59 DUPLICATE REMOVE L2 (32 DUPLICATES REMOVED) => d 13 1-10 ANSWER 1 OF 59 CAPLUS COPYRIGHT 2004 ACS on STN 2004:174715 CAPLUS Matrix-Assisted Laser Desorption Ionization Time-of-Flight Mass Spectrometry Analysis of ***Zeins*** in Mature Maize Kernels Adams, Whitney R.; Huang, Shihshieh; Kriz, Alan L.; Luethy, Michael H. Mystic Research, Monsanto Company, Mystic, CT, 06355, USA Journal of Agricultural and Food Chemistry (2004), 52(7), 1842-1849 CODEN: JAFCAU; ISSN: 0021-8561 American Chemical Society Journal English RE.CNT 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 2 OF 59 CAPLUS COPYRIGHT 2004 ACS on STN 2003:242097 CAPLUS DN 138:267201 Pesticidal compositions for coating plant propagation material containing anthranilamides Berger, Richard Alan; Flexner, John Lindsey E. I. Du Pont de Nemours & Co., USA PCT Int. Appl., 147 pp. CODEN: PIXXD2

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DUPLICATE 11

ACCESSION NUMBER: 97:33751 AGRICOLA

DOCUMENT NUMBER: IND20564663

TITLE: The maize ***gamma*** - ***zein*** sequesters

alpha- ***zein*** and stabilizes its accumulation in ***protein*** bodies of transgenic tobacco

endosperm.

AUTHOR(S): Coleman, C.E.; Herman, E.M.; Takasaki, K.; Larkins,

B.A.

CORPORATE SOURCE: Brigham Young University, Provo, UT.

AVAILABILITY: DNAL (OK725.P532)

SOURCE: The Plant cell, Dec 1996. Vol. 8, No. 12. p. 2335-2345

Publisher: [Rockville, MD : American Society of Plant

Physiologists, c1989-

CODEN: PLCEEW; ISSN: 1040-4651

NOTE: Includes references
PUB. COUNTRY: Maryland; United States

DOCUMENT TYPE: Article

FILE SEGMENT: U.S. Imprints not USDA, Experiment or Extension

LANGUAGE: English

Zeins are ***seed*** storage ***proteins*** that form accretions called ***protein*** bodies in the rough endoplasmic reticulum of maize endosperm cells. Four types of ***zeins***, alpha, beta, ***gamma***, and delta, aggregate in a distinctive spatial pattern within the ***protein*** body. We created transgenic tobacco plants expressing alpha- ***zein***, ***gamma*** - ***zein***, or both to examine the interactions between these ***proteins*** leading to the formation of ***protein*** bodies in the endosperm. Whereas ***gamma*** - ***zein*** accumulated in ***seeds*** of these plants, stable accumulation of alpha- ***zein*** required simultaneous synthesis of ***gamma*** - ***zein***. The ***zein*** ***proteins*** formed accretions in the endoplasmic reticulum similar

those in maize endosperm. ***Protein*** bodies were also found in
protein storage vacuoles. The accumulation of both types of
zeins peaked early in development and declined during maturation.
Even in the presence of ***gamma*** - ***zein*** , there was a
turnover of alpha- ***zein*** , suggesting that the interaction between
the two ***proteins*** might be transitory. We suggest that
gamma - ***zein*** plays an important role in ***protein***
body formation and demonstrate the utility of tobacco for studying
interactions between different ***zein*** .

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L4 35 GAMMA(W) ZEIN AND TRANSFORM?

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- L5 ANSWER 1 OF 20 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Production of peptides and proteins by accumulation in plant endoplasmic reticulum-derived protein bodies
- L5 ANSWER 2 OF 20 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Self-processing transgenic plants and plant parts expressing hyperthermophilic processing enzymes
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 (2004) on STN DUPLICATE 1
- TI Expression of the ***gamma*** ***zein*** protein of maize in seeds of transgenic barley: effects on grain composition and properties.
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 DUPLICATE 2
- TI Zein accumulation in forage species (Lotus corniculatus and Medicago sativa) and co-expression of the ***gamma*** ***zein*** :KDEL and beta-zein: KDEL polypeptides in tobacco leaf.
- L5 ANSWER 5 OF 20 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN DUPLICATE 3
- TI Combination of viral promoter sequences to generate highly active promoters for heterologous therapeutic protein over-expression in plants.
- ANSWER 6 OF 20 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2004) on STN
- TI Expression of a synthetic E. coli heat-labile enterotoxin B sub-unit (LT-B) in maize.
- L5 ANSWER 7 OF 20 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN DUPLICATE 4 $\,$
- TI Expression of a synthetic porcine alpha-lactalbumin gene in the kernels of transgenic maize.
- L5 ANSWER 8 OF 20 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Geminivirus replicases and the genes encoding them and their use to create polyploid plant cells
- L5 ANSWER 9 OF 20 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Methods of using viral replicase polynucleotides and polypeptides in transgenic plants
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| <u>L7</u> | zein and lysine and enriched | 97 | <u>L7</u> |
| <u>L6</u> | maize and increased adj lysine | 44 | <u>L6</u> |
| <u>L,5</u> | maize and increased adj lusine | 0 | <u>L5</u> |
| DB=DWF | PI; PLUR=YES; OP=OR | | |
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| L2 | WO adj 9315221 | 4 | L2 |
| <u>L_1</u> | WO 9315221 | 915453 | <u>L1</u> |

END OF SEARCH HISTORY